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10/719,976	11/21/2003	Xuedong Song	KCX-693 (19341)	1744
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)		
	10/719,976	SONG, XUEDONG		
Office Action Summary	Examiner	Art Unit		
	Jacqueline DiRamio	1641		
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the o	correspondence address		
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA  - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication.  - If NO period for reply is specified above, the maximum statutory period verailure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tir vill apply and will expire SIX (6) MONTHS from a cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).		
Status				
Responsive to communication(s) filed on 12 M     This action is <b>FINAL</b> . 2b) ☑ This     Since this application is in condition for alloware closed in accordance with the practice under E	action is non-final.			
Disposition of Claims				
4) ☐ Claim(s) 1-6, 9-13, and 39-45 is/are pending in 4a) Of the above claim(s) is/are withdraw 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) 1-6,9-13 and 39-45 is/are rejected. 7) ☐ Claim(s) 1 and 12 is/are objected to. 8) ☐ Claim(s) are subject to restriction and/or	wn from consideration.			
Application Papers				
9) ☐ The specification is objected to by the Examine 10) ☑ The drawing(s) filed on 26 April 2004 is/are: a) Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) ☐ The oath or declaration is objected to by the Ex	☑ accepted or b)☐ objected to drawing(s) be held in abeyance. Section is required if the drawing(s) is ob	e 37 CFR 1.85(a). jected to. See 37 CFR 1.121(d).		
Priority under 35 U.S.C. § 119				
<ul> <li>12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).</li> <li>a) All b) Some * c) None of:</li> <li>1. Certified copies of the priority documents have been received.</li> <li>2. Certified copies of the priority documents have been received in Application No.</li> <li>3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).</li> <li>* See the attached detailed Office action for a list of the certified copies not received.</li> </ul>				
Attachment(s)  1) Notice of References Cited (PTO-892)  2) Notice of Draftsperson's Patent Drawing Review (PTO-948)  3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)  Paper No(s)/Mail Date	4)  Interview Summary Paper No(s)/Mail Di 5)  Notice of Informal F 6)  Other:			

## **DETAILED ACTION**

### Continued Examination Under 37 CFR 1.114

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on May 12, 2008 has been entered.

#### Status of the Claims

Applicant's amendments to claims 1, 9, and 11 are acknowledged, as well as the cancellation of claims 7, 8, and 14 - 38, and the addition of new claims 39 - 45.

Currently, claims 1 - 6, 9 - 13, and 39 - 45 are pending and under examination.

## Withdrawn Rejections

All previous rejections of the claims under 35 U.S.C. 102 and 103 are withdrawn in view of Applicant's amendments and arguments filed May 12, 2008.

## Response to Arguments

Applicant's arguments, see p6-8, filed May 12, 2008, with respect to the rejection(s) of the claim(s) under 35 U.S.C. 103(a) as being unpatentable over Daniels et al. (US 2006/0008921) in view of Jou et al. (US 5,670,381) have been fully

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considered and are persuasive. Applicant's argument that neither Daniels et al. nor Jou et al. teach the amendment to claim 1 requiring the "calibration zone" to be "positioned between said detection zone and said compensation zone" is found persuasive.

Therefore, the rejections have been withdrawn. However, upon further consideration, a

#### **NEW GROUNDS OF REJECTION**

# Claim Objections

Claims 1 and 12 are objected to because of the following informalities:

Claims 1 and 12 recite the term "third capture reagent," however, because there is no longer a "second capture reagent," this term is incorrect and confusing.

Appropriate correction is required.

new ground(s) of rejection is made and presented below.

## Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

The factual inquiries set forth in *Graham* **v.** *John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

1. Determining the scope and contents of the prior art.

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2. Ascertaining the differences between the prior art and the claims at issue.

- 3. Resolving the level of ordinary skill in the pertinent art.
- 4. Considering objective evidence present in the application indicating obviousness or nonobviousness.

Claims 1 - 6, 9 - 13, and 39 - 45 are rejected under 35 U.S.C. 103(a) as being unpatentable over Rylatt et al. (WO 97/09620), as evidenced by Fitzpatrick et al. (US 6,121,008), in view of Jou et al. (US 5,670,381).

Rylatt et al. teach a flow-through assay device for detecting the presence or quantity of an analyte residing in a test sample, said flow-through assay device comprising a porous membrane 207, said porous membrane being in communication with analyte detection agents 208 (probes) and calibration agents 209 (probes), said analyte detection agents being conjugated with a specific binding partner (member) for the analyte, said porous membrane defining:

a test (detection) zone 204 within which is immobilized an analyte receptor 215 (first capture reagent), said analyte receptor being configured to bind to at least a portion of said complexes formed between the analyte and said conjugated analyte detection agents to generate a test (detection) signal having an intensity;

a procedural control (compensation) zone 212 located downstream from said test zone, wherein a binding agent is immobilized within said procedural control zone, said binding agent configured to bind to said conjugated analyte detection agents and complexes formed between the analyte and said conjugated analyte detection agents passing through said test zone to generate a control (compensation) signal having an intensity; and

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a calibration zone(s) 211 within which a calibration agent receptor 214 (third capture reagent) is immobilized, said calibration agent receptor being configured to bind to said calibration agents to generate a calibration signal that is substantially constant relative to the intensities of said test signal and said control signal, said calibration zone being positioned between said test zone and said procedural control zone:

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wherein the amount of analyte within the test sample is proportional to the ratio of said test signal intensity to said control signal intensity, as calibrated by said calibration signal intensity (see Figure 2a; p7, lines 25-29; p8, lines 20-29; p10, lines 27-30; p12, lines 25-30; p14, lines 2-19; p15, lines 11-12; and Example 1).

With respect to the limitation of Applicant's claim 1 reciting "wherein the intensity of said compensation signal is inversely proportional to the intensity of said detection signal," the signal generated in the control zone of Rylatt et al. would create a signal that is inversely proportional to the intensity of the test signal because the control zone is binding to the unbound conjugated detectable labels, wherein the amount of unbound detectable labels would be inversely proportional to the amount of analyte in the test sample. Since the test zone is binding to the complexes created between the analyte and the detectable label, the test signal would be proportional to the amount of analyte in the sample. Therefore, the control signal created in the control zone of Rylatt et al. would in fact be inversely proportional to the test signal created in the test zone (see Fitzpatrick et al.: column 2, lines 19-67; and column 3, lines 1-9).

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However, Rylatt et al. fail to teach that the binding agent immobilized in the procedural control zone (compensation zone) comprises a polyelectrolyte, wherein said polyelectrolyte has a net charge opposite to that of the detection agents (probes).

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Jou et al. teach a device for performing an assay comprising a porous material containing a detection zone with an immobilized capture reagent. The device utilizes a diffusive indicator reagent, which comprises a specific binding member attached to a detectable label, and a diffusive capture reagent, which comprises a charged substance, wherein the reagents are contacted with an analyte of interest to form a complex. The complex binds to the immobilized capture reagent in the detection zone through ion-capture, wherein the capture reagent is oppositely charged with respect to the charged substance of the analyte complex. The immobilized capture reagent preferably comprises an anionic or cationic polymeric substance (polyelectrolyte), which enables the production of a generic solid phase device for use in specific binding assays. Assay procedures for many different analytes can use the same solid phase material which contains a predetermined zone of anionic or cationic capture polymer rather than an immobilized binding member capable of binding only a specific analyte as found in conventional flow-through or test-strip devices. Further, the ion-capture technique increases the potential number of complexes that can be immobilized on the solid support (see column 6, lines 25-40; column 7, lines 1-46; column 10, lines 63-65; column 19, lines 29-67; column 24, lines 49-67; column 25, lines 1-57; and column 26, lines 18-61).

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Therefore, it would have been obvious to one of ordinary skill in the art at the time the invention was made to modify the procedural control zone of the device of Rylatt et al. to comprise a polyelectrolyte that is oppositely charged with respect to a charged substance of a diffusive indicator reagent as taught by Jou et al. because Jou et al. teach the benefit of using an anionic or cationic polymeric substance as the immobilized capture reagent in a detection zone because the polymeric substance allows for the binding of a charged conjugated substance or complex to a solid phase support material through ion-capture, which increases the potential number of complexes that can be immobilized on the solid support and allows for the production of a generic solid phase device, wherein many different analytes can use the same solid phase material which contains a predetermined zone of anionic or cationic capture polymer rather than an immobilized binding member capable of binding only a specific analyte as found in conventional flow-through or test-strip devices.

With respect to Applicant's claims 2-4 and 42, Rylatt et al. teach that the analyte detection agents can comprise a fluorophore (luminescent compound), a chemiluminescent molecule, a chromogen, a radioisotope, or a direct visual label, such as a colloidal metal or a latex particle (see p8, lines 22-29).

With respect to Applicant's claims 5 and 44, Rylatt et al. teach that the specific binding partner is preferably an antibody (see Example 1).

With respect to Applicant's claims 6 and 45, Rylatt et al. teach that the analyte receptor (first capture reagent) preferably comprises an antibody (see Example 1).

With respect to Applicant's claims 9 and 11, Jou et al. teach that the charged polymeric substance (polyelectrolyte) can be anionic or cationic (see column 7, lines 1-46; and column 19, lines 29-36).

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With respect to Applicant's claim 10, Jou et al. teach various charged polymers for the polyelectrolyte, which include combinations of those recited in Applicant's claim 10 (see column 19, lines 29-67; and column 20, lines 1-67).

With respect to Applicant's claim 12, Rylatt et al. teach that the calibration agent receptor (third capture reagent) and calibration agent can comprise streptavidin and biotin (see Example 1).

With respect to Applicant's claim 13, Rylatt et al. teach that the device can comprise a sandwich-type assay device (see Example 1).

With respect to Applicant's claims 39 – 41, both Rylatt et al. and Jou et al. teach the direct or indirect and covalent or non-covalent attachment of reagents, or polyelectrolyte, to the surface of the porous membrane, as well as the use of spacer molecules (functional groups) to bind the receptor reagents to the porous membrane (see p11, lines 12-18 of Rylatt et al; and column 28, lines 1-13 of Jou et al).

With respect to Applicant's claim 43, the combination of Rylatt et al. in view of Jou et al. would result in the binding agent of the procedural control (compensation) zone of Rylatt et al. being substituted with the polyelectrolyte of Jou et al., which would result in the procedural control zone being free of biological capture reagents.

## Conclusion

No claims are allowed.

The following prior art made of record and not relied upon is considered pertinent to applicant's disclosure:

Inganas et al. (US 2006/0175193) teach a complex created between a polyelectrolyte and one or more receptor molecules specific for a target biomolecule analyte for use as a probe in biomolecular interactions (see Abstract; and paragraphs [0006] and [0022]).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jacqueline DiRamio whose telephone number is 571-272-8785. The examiner can normally be reached on M-F 9-5:30.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Long Le can be reached on 571-272-0823. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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/Jacqueline DiRamio/ Examiner, Art Unit 1641

> /Mark L. Shibuya, Ph.D./ Supervisory Patent Examiner, Art Unit 1641